

United States Patent and Trademark Office

UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER OF PATENTS AND TRADEMARKS Washington, D.C. 20231 www.uspto.gov

CONFIRMATION NO. APPLICATION NO. FILING DATE FIRST NAMED INVENTOR ATTORNEY DOCKET NO. PF-0420-2 DIV 8687 09/757,716 01/09/2001 Holly Magna 7590 02/14/2002 INCYTE GENOMICS, INC. **EXAMINER** PATENT DEPARTMENT SAIDHA, TEKCHAND 3160 Porter Drive Palo Alto, CA 94304 ART UNIT PAPER NUMBER 1652 DATE MAILED: 02/14/2002

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)			
	09/757,716	MAGNA ET AL.			
Office Action Summary	Examiner	Art Unit			
	Tekchand Saidha	1652			
The MAILING DATE of this communication app Period for Reply	ars on the cover shet with the	he corr spond nce address			
A SHORTENED STATUTORY PERIOD FOR REPLY THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a reply - If NO period for reply is specified above, the maximum statutory period v - Failure to reply within the set or extended period for reply will, by statute - Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b). Status	66(a). In no event, however, may a reply be within the statutory minimum of thirty (30) will apply and will expire SIX (6) MONTHS cause the application to become ABANDO	oe timely filed) days will be considered timely. from the mailing date of this communication. ONED (35 U.S.C. § 133).			
1) Responsive to communication(s) filed on <u>01 f</u>	ebruary, 2002 .				
2a)☐ This action is FINAL . 2b)⊠ Th	s action is non-final.				
3) Since this application is in condition for allowated closed in accordance with the practice under	•	• •			
Disposition of Claims					
4)⊠ Claim(s) 1,10,14-16 and 27-44 is/are pending in the application.					
4a) Of the above claim(s) is/are withdraw	vn from consideration.				
5) Claim(s) is/are allowed.					
6)☐ Claim(s) is/are rejected.					
7) Claim(s) is/are objected to.					
8)⊠ Claim(s) <u>1,10,14-16 and 27-44</u> are subject to r	estriction and/or election requ	irement.			
Application Papers					
9)☐ The specification is objected to by the Examine	r.				
10)☐ The drawing(s) filed on is/are: a)☐ accept	ted or b) objected to by the E	Examiner.			
Applicant may not request that any objection to the	e drawing(s) be held in abeyance	e. See 37 CFR 1.85(a).			
11) The proposed drawing correction filed on	is: a)☐ approved b)☐ disap	oproved by the Examiner.			
If approved, corrected drawings are required in rep	ly to this Office action.				
12)☐ The oath or declaration is objected to by the Ex	aminer.				
Priority under 35 U.S.C. §§ 119 and 120					
13) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).					
a) ☐ All b) ☐ Some * c) ☐ None of:					
1. Certified copies of the priority documents have been received.					
2. Certified copies of the priority documents have been received in Application No					
3. Copies of the certified copies of the prior application from the International Bu * See the attached detailed Office action for a list	reau (PCT Rule 17.2(a)).				
14) ☐ Acknowledgment is made of a claim for domesti					
a) The translation of the foreign language pro	visional application has been	received.			
15) Acknowledgment is made of a claim for domestic Attachment(s)	c priority under 35 U.S.C. 99	120 anu/01 121.			
1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO-1449) Paper No(s)	5) Notice of Inform	mary (PTO-413) Paper No(s) mal Patent Application (PTO-152)			

Art Unit: 1652

Restriction to one of the following inventions is required under 35 U.S.C. 121:

- I. Claim 1, drawn to human nucleotide pyrophosphohydrolase-2 (NTPPH-2), classified in class 435, subclass 195.
- II. Claims 10 & 29-44, drawn to antibody, composition comprising the antibody and methods of use, classified in class 424, subclass 130.1.
- III. Claims 14-16 & 28 drawn to a method of detecting a polynucleotide, classified in class 435, subclass 6.
- IV. Claim 27, drawn to a method of screening compounds for altered expression of target polynucleotide, classified in class 435, subclass 69.2.

The inventions are distinct, each from the other because of the following reasons:

Inventions of group I and group II are distinct because protein and antibody are chemically and biologically distinct molecules. Antibody and protein have fundamentally different molecular structure, each with its own set of functionality. Antibodies, for example are formed in the B-cells and are useful for binding to particular residues. Proteins do not function to bind in the particular immunological way that antibodies do, and therefore have different specificities for different substrates, and do not purport to have the kinds of specific activity that antibodies have.

3. Inventions of group I and group III are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (M.P.E.P. § 806.05(h)). The protein of Group I can be used in a variety of ways

Application/Control Number: 09/757,716

Art Unit: 1652

other than for method of hybridization by the invention of Group III. The protein can be used to find other inhibitors/activators that can be used in the treatment or diagnoses of certain illnesses. At the minimum the protein can be used to delineate molecular weight parameters in a protein gel electrophoresis assay. The inventions of group I and IV are also different for similar reasons i.e., the protein can used for the enzymatic formation of the product rather than in the method of assessing toxicity.

- 4. The Inventions of group II and group IV are independent and distinct because the antibody and a screening method to identify a compound are functionally distinct and neither requires the other to practice the invention.
- 5. Inventions II and III are patentably distinct from each other. The antibodies against the NTPPH-2 enzyme of Group II and the hybridization method requiring polynucleotide of Group III do not require each other for their practice; have separate utilities, structure and function. Invention of Group III and IV though use the same polynucleotide but are distinct because of separate utilities and distinct method steps. These groups have acquired separate status in the art and separate fields of search as further evidenced by their separate classification.

Because these inventions are distinct for the reasons given above and have acquired a separate status in the art as shown by their different classification, restriction for examination purposes as indicated is proper.

A telephone call was made to Richard Ekstrom/Margerrette on 2.1.02 to request an oral election to the above restriction requirement, but did not result in an election being made.



Creation date: 09-08-2003

Indexing Officer: NDUBOSE - NEFERTITI DUBOSE

Team: OIPEBackFileIndexing

Dossier: 09757716

Legal Date: 03-25-2002

No.	Doccode	Number of pages
1	A	2
2	CLM	4
3	REM	2

Total number of pages: 8		
Total Hambel of pages. o		
Remarks:		

Order of re-scan issued on